AMENDMENT TO THE CLAIMS

Please replace all prior versions and listings of claims with the amended claims as follows:

1. (Currently amended) A compound of formula I:

or a pharmaceutically acceptable salt thereof, wherein:

Ht is pyrrol-3-yl having R³ and QR⁴ substituents;

A-B is N-O or O-N;

R1 is hydrogen or -NHR;

T is a valence bond;

Q is -C(O) or $-SO_2$ -;

each R is independently selected from hydrogen or an optionally substituted aliphatic group having one to six carbons;

R² is an aryl group substituted with up to three R⁸ substituents;

R³ is hydrogen;

R⁴ is -R⁶ or -NHR⁶;

each R⁶ is independently selected from -(CH₂)_yR⁷;

y is 0-6;

each R⁷ is an optionally substituted group independently selected from aryl, heteroaryl, or heterocyclyl;

each R^8 is independently selected from haloger, -R', or -OR'; wherein each R' is independently selected from hydrogen[[,]] or an optionally substituted C_{1-12} aliphatic [[; and]].

2. (Currently amended) The compound according to claim 1 having the formula:

or a pharmaceutically acceptable salt thereof, wherein the variables Ht, T, R¹, and R² are as defined in claim 1.

3. (Currently amended) The compound according to claim 2 having the formula:

$$\begin{array}{c|c}
H \\
N \\
Q-R^4
\end{array}$$

$$\begin{array}{c|c}
T-R^2
\end{array}$$

II.A

or a pharmaceutically acceptable salt thereof, wherein the variables Q, T, R¹, R², R³, and R⁴ are as defined in claim-1.

- 4. (Previously amended) The compound according to claim 3, wherein said compound has one or more features selected from the group consisting of:
 - (a) Q is -CO-;
 - (b) R1 is hydrogen; and
 - (c) R⁷ is an optionally substituted heterocyclyl group.
 - 5. (Previously amended) The compound according to claim 4, wherein:
 - (a) Q is -CO-;
 - (b) R¹ is hydrogen; and
 - (c) R⁷ is an optionally substituted heterocyclyl group.

6. (Currently amended) The compound according to claim 1 having the formula:

$$O_{R^1} \xrightarrow{Ht} T - R^2$$

III

or a pharmaceutically acceptable salt thereof, wherein the variables Ht, T, R^1 , and R^2 are as defined in claim 1.

7. (Currently amended) The compound according to claim 6 having the formula:

III-A

or a pharmaceutically acceptable salt thereof, v/herein the variables $Q, T, R^{4}, R^{2}, R^{3}$, and R^{4} are as defined in claim 1.

- 8. (Previously amended) The compound according to claim 7, wherein said compound has one or more features selected from the group consisting of:
 - (a) Q is -CO-;
 - (b) R1 is hydrogen; and
 - (c) R⁷ is an optionally substituted heterocyclyl group.
 - 9. (Previously amended) The compound according to claim 8, wherein:
 - (a) Q is -CO-;
 - (b) R1 is hydrogen; and
 - (c) R⁷ is an optionally substituted heterocyclyl group.

10. (Currently amended) The compound according to claim 1 having the formula:

or a pharmaceutically acceptable salt thereof, wherein the variables Ht, T, R¹, and R² are as defined in claim 1.

11. (Currently amended) The compound according to claim 10 having the formula:

TV.A

or a pharmaceutically acceptable salt thereof, wherein the variables Q, T, R^1 , R^2 , R^3 , and R^4 are as defined in claim 1.

- 12. (Previously amended) The compound according to claim 11, wherein said compound has one or more features selected from the group consisting of:
 - (a) Q is -CO-; and
 - (b) R⁷ is an optionally substituted heterocyclyl group.
 - 13. (Previously amended) The compound according to claim 12, wherein:
 - (a) Q is -CO-; and
 - (b) R⁷ is an optionally substituted heterocyclyl group.
 - 14-17. (Canceled)

18. (Currently amended) The compound according to claim 1, wherein said compound is selected from the following Table 1 and Table 2 compounds having formulae II-A and IV-A:

$$\begin{array}{c}
H \\
N \\
-Q-R^4
\end{array}$$

$$\begin{array}{c}
T-R^2
\end{array}$$

Compounds of Formula II-A

No.	T-R ²	Q-R ⁴	
ПА-2	2-chlorophenyl	CONHCH₂(Ph)	
ПА-3	2-chlorophenyl	CO(morpholin-4-yl)	
ПА-4	4-methoxyphenyl	CONHCH2(pyridin-4-yl)	
ПА-5	3-fluorophenyl	CONHCH₂(pyridin-4-yl)	
ПА-6	3-methoxyphenyl	CONHCH₂(pyridin-4-yl)	
ПА-7	2,5-dimethoxyphenyl	CONHCH2(pyridin-4-yl)	
ПА-8	3,4-difluorophenyl	CONHCH₂(pyridin-4-yl)	
ПА-9	2,3-difluorophenyl	CONHCH₂(pyridin-4-yl)	
IIA-10	2,5-difluorophenyl	CONHCH ₂ (pyridin-4-yl)	
ПА-11	4-methoxyphenyl	CONHCH ₂ (pyridin-3-yl)	
ПА-12	3-fluorophenyl	CONHCH ₂ (pyridin-3-yl)	
IIA-13	3-methoxyphenyl	CONHCH ₂ (pyridin-3-yl)	
ПА-14	2,5-dimethoxyphenyl	CONHCH ₂ (pyridin-3-yl)	
IIA-15	3,4-difluorophenyl	CONHCH2(pyridin-3-yl)	
ПА-16	2,3-difluorophenyl	CONHCH₂(pyridin-3-yl)	
IIA-17	2,5-difluorophenyl	CONHCH2(pyridin-3-yl)	
ПА-18	4-methoxyphenyl	CONHCH ₂ (tetrahydrofuran-2-yl)	
IIA-19	3-fluorophenyl	CONHCH2(tetrahydrofuran-2-yl)	
ПА-20	3-methoxyphenyl	CONHCH2(tetrahydrofuran-2-yl)	
ПА-21	2,5-dimethoxyphenyl	CONHCH ₂ (tetrahydrofuran-2-yl)	
ПА-22	3,4-difluorophenyl	CONHCH2(tetrahydrofuran-2-yl)	
IIA-23	2,3-difluorophenyl	CONHCH2(tetrahydrofuran-2-yl)	
ПА-24	2,5-difluorophenyl	CONHCH2(tetrahydrofuran-2-yl)	
IIA-25	4-fluorophenyl	CONHCH₂(1-Et-pyrrolidin-2-yl)	

No.	No. $T-R^2$ $Q-R^4$			
IIA-26	4-methoxyphenyl	CONHCH2(1-Et-pyrrolidin-2-yl)		
IIA-27	3-fluorophenyl	CONHCH2(1-Et-pyrrolidin-2-yl)		
IIA-28	3-methoxyphenyl	CONHCH2(1-Et-pyrrolidin-2-yl)		
IIA-29	2,5-dimethoxyphenyl	CONHCH2(1-Et-pyrrolidin-2-yl)		
ПА-30	3,4-difluorophenyl	CONHCH2(1-Et-pyrrolidin-2-yl)		
IIA-31	2,3-difluorophenyl	CONHCH2(1-Et-pyrrolidin-2-yl)		
IIA-32	2,5-difluorophenyl	CONHCH₂(1-Et-pyrrolidin-2-yl)		
IIA-33	4-fluorophenyl	CO(morpholin-4-yl)		
ПА-34	4-methoxyphenyl	CO(morpholin-4-yl)		
IIA-35	3-fluorophenyl	CO(morpholin-4-yl)		
IIA-36	3-methoxyphenyl	CO(morpholin-4-yl)		
IIA-37	2,5-dimethoxyphenyl	CO(morpholin-4-yl)		
IIA-38	2,3-difluorophenyl	CO(morpholin-4-yl)		
IIA-39	2,5-difluorophenyl	CO(morpholin-4-yl)		
IIA-40	4-fluorophenyl	CO(4-Me-piperazin-1-yl)		
IIA-41	4-methoxyphenyl	CO(4-Me-piperazin-1-yl)		
ПА-42	3-fluorophenyl	CO(4-Me-piperazin-1-yl)		
ПА-43	3-methoxyphenyl	CO(4-Me-piperazin-1-yl)		
IIA-44	2,5-dimethoxyphenyl	CO(4-Me-piperazin-1-yl)		
IIA-45	2,3-difluorophenyl	CO(4-Me-piperazin-1-yl)		
IIA-46	2,5-difluorophenyl	CO(4-Me-piperazin-1-yl)		
IIA-47	3-chlorophenyl	CONHCH ₂ (pyridin-4-yl)		
IIA-48	3-chlorophenyl	CONHCH ₂ (pyridin-3-yl)		
ПА-49	3-chlorophenyl	CONHCH2(tetrahydrofuran-2-yl)		
IIA-50	3-chlorophenyl	CONHCH ₂ (1-Et-pyrrolidin-2-yl)		
ПА-51	3-chlorophenyl	CO(4-Me-piperazin-1-yl)		
IIA-52	4-chlorophenyl	CONHCH ₂ (pyridin-4-yl)		
ПА-53	4-chlorophenyl	CONHCH ₂ (pyridin-3-yl)		
IIA-54	4-chlorophenyl	CONHCH2(tetrahydrofuran-2-yl)		
IIA-55	4-chlorophenyl	CONHCH2(1-Et-pyrrolidin-2-yl)		
IIA-56	4-chlorophenyl	CO(morpholin-4-yl)		
IIA-57	4-chlorophenyl	CO(4-Me-piperazin-1-yl)		
ПА-58	3,4-dichlorophenyl	CONHCH₂(pyridin-3-yl)		
IIA-59	3,4-dichlorophenyl	CONHCH2(1-Et-pyrrolidin-2-yl)		

Т				
No.	T-R ²	Q-R ⁴		
IIA-60	3,4-dichlorophenyl	CO(morpholin-4-yl)		
IIA-61	3,4-dichlorophenyl	CO(4-Me-piperazin-l-yl)		
IIA-62	2-F-3-chlorophenyl	CONHCH₂(pyridin-4-yl)		
IIA-63	2-F-3-chlorophenyl	CONHCH₂(pyridin-3-yl)		
IIA-64	2-F-3-chlorophenyl	CONHCH ₂ (tetrahydrofuran-2-yl)		
IIA-65	2-F-3-chlorophenyl	CONHCH2(1-Et-pyrrolidin-2-yl)		
IIA-66	2-F-3-chlorophenyl	CO(morpholin-4-yl)		
ПА-67	2-F-3-chlorophenyl	CO(4-Me-piperazin-1-yl)		
ДА-68	3-Cl-4-fluorophenyl	CONHCH2(pyridin-4-yl)		
IIA-69	3-Cl-4-fluorophenyl	CONHCH ₂ (pyridin-3-yl)		
ПА-70	3-Cl-4-fluorophenyl	CONHCH2(tetrahydrofuran-2-yl)		
IIA-71	3-Cl-4-fluorophenyl	CONHCH ₂ (1-Et-pyrrolidin-2-yl)		
ПА-72	3-Cl-4-fluorophenyl	CO(morpholin-4-yl)		
ILA-73	3-Cl-4-fluorophenyl	CO(4-Me-piperazin-1-yl)		
11A-74	3,4-dimethoxyphenyl	CONHCH2(pyridin-4-yl)		
ПА-75	3,4-dimethoxyphenyl	CONHCH ₂ (pyridin-3-yl)		
IIA-76	3,4-dimethoxyphenyl	CONHCH ₂ (tetrahydrofuran-2-yl)		
ПА-77	3,4-dimethoxyphenyl	CONHCH ₂ (1-Et-pyrrolidin-2-yl)		
IIA-78	3,4-dimethoxyphenyl	CO(morpholin-4-yl)		
ПА-79	3,4-dimethoxyphenyl	CO(4-Me-piperazin-1-yl)		
IIA-80	4-benzo[1,3]dioxol-5-yl	CONHCH2(pyridin-4-yl)		
IIA-81	4-benzo[1,3]dioxol-5-yl	CONHCH2(pyridin-3-yl)		
IIA-82	4-benzo[1,3]dioxol-5-yl	CONHCH ₂ (tetrahydrofuran-2-yl)		
IIA-83	4-benzo[1,3]dioxol-5-yl	CONHCH2(1-Et-pyrrolidin-2-yl)		
IIA-84	4-benzo[1,3]dioxol-5-yl	CO(morpholin-4-yl)		
IIA-85	4-benzo[1,3]dioxol-5-yl	CO(4-Me-piperazin-1-yl)		
ПА-86	3,5-dichlorophenyl	CONHCH₂(pyridin-4-yl)		
IIA-87	3,5-dichlorophenyl	CONHCH₂(pyridin-3-yl)		
IIA-88	3,5-dichlorophenyl	CONHCH ₂ (tetrahydrofuran-2-yl)		
IIA-89	3,5-dichlorophenyl	CONHCH2(1-Et-pyrrolidin-2-yl)		
IIA-90	3,5-dichlorophenyl	CO(morpholin-4-yl)		
IIA-91	3,5-dichlorophenyl	CO(4-Me-piperazin-1-yl)		
IIA 92	3 Cl 4 SO ₂ NH ₂ phenyl	CO(morpholin 4-yl)		
IIA-93	3-chlorophenyl	CO(morpholin-4-yl)		

No.	T-R ²	Q-R ⁴		
IIA-94	phenyl	pyridin 4-yl		
IIA 95	2-chlorophenyl	morpholin 4-yl		
IIA 96	2 chlorophenyl	CH ₂ (morpholin 4 yl)		
IIA-97	4 methoxyphenyl	СН _{«(ругіdin 4-yl)}		
IIA-106	phenyl .	3 ¹ N N		
IIA-107	phenyl	3 ¹ N		
IIA-108	3,4-dimethoxyphenyl	32 N N F		
IIA-109	3-chlorophenyl	3 ² N N		
ПА-110	3-chlorophenyl	3 ¹ N		
ПА-111	3-methylphenyl	ئ ^ا ين		
ПА-114	2-fluoro-3-chlorophenyl	ئ ^ا ر ال		
ПА-115	3-chlorophenyl	ş ^N N OH3		

No.	T-R ²	Q-R ⁴		
ПА-116	3,4-dimethoxyphenyl	ş ^l N		
НА -117	3,4-dimethoxyphenyl	2 N OH		
IIA-119	3-methylphenyl	2 NOH		
ПА-120	2-fluoro-3-chlorophenyl	z ^l n ,		
ПА -121	2-fluoro-3-chlorophenyl	Syll N N N N Me		
ПĄ-122	2-fluoro-3-chlorophenyl	3 ² N N N N N N N N N N N N N N N N N N N		
ПА-123	3-chlorophenyl	ئ ^ا ا		
ПА-124	3,4-dimethoxyphenyl	2 ¹ N		
IIA-125	2-fluoro-3-chlorophenyi	2 ¹ 2 5 6 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7		
ПА-126	2-fluoro-3-chlorophenyl	3 ^N NNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN		

No.	T-R ²	Q-R ⁴		
ПА-130	phenyl	2 ^N NNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN		
ПА-131	phenyl	N N N N N N N N N N N N N N N N N N N		
ПА-132	phenyl	ئ ^ا »		
IIA-133	phenyl	², ¹ , N,		
IIA-134	phenyl	Ž ^N N⊃ C		
IIA-135	3,4-dimethoxyphenyl	12 CO T T CO		
IIA-136	3,4-dimethoxyphenyl	ئ ^ا ين		
ПА-137	3,4-dimethoxyphenyl	ş ^l n⊃n		
ПА-138	3-methylphenyl	3100		
IIA-139	3-methylphenyl	2 N N		
ПА-140	3-methylphenyl	3 ¹ N		

No.	T-R ²	Q-R ⁴	
ПА-141	2-fluoro_[[,]]3-chlorophenyl	2 ¹ N	
IIA-142	3-chlorophenyl		
ПА-143	3-chlorophenyl		
IIA-144	3-chlorophenyl	2 ¹ N	
ПА-145	3-chlorophenyl	ş ^Î N C	
ПА-146	3-chlorophenyl	3 ¹ NNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN	
IIA-148	phenyl	y N N N N N N N N N N N N N N N N N N N	
IIA-150	3,4-dimethoxyphenyl	ب ^ا ا	
ПА-151	3-methylphenyl	3 ¹ 2 1	
11A-152	3-methylphenyl	Z-N CH₃	

No.	T-R ²	Q-R ⁴		
ПА-153	phenyl	2 NOH		
ПА-154	phenyl	Š CH3		
HA -155	phenyl			
IIA-156	3,4-dimethoxyphenyl	Z N N N OME		
IIA-157	3,4-dimethoxyphenyl	² N N CH ₀		
IIA -159	3-methylphenyl	ş.Å _N → OH		
ПА-160	3-chlorophenyl	3 N OH		
IIA-161	phenyl	چگ _{ای} ک		
IIA-162	3-chlorophenyl	, ² , ³		
ПА-163	3,4-dimethoxyphenyl	δ _N CH ³		

No.	T-R ²	Q-R⁴		
IIA-164	3-chlorophenyl	², ll N CH₃		
IIA-165	phenyl	ž, N OH		
IIA-167	pbenyl	Ž N CH³		
IIA-168	3,4-dimethoxyphenyl	2 NOH		
ПА-169	3,4-dimethoxyphenyl	3 ¹ 2		
IIA-170	3,4-dimethoxyphenyl	ş ^l N CN C		
HA -171	3-methylphenyl	Z, N, OH		
ПА-172	3-methylphenyl	ş l N N N N N N N N N N N N N N N N N N		
ЦА -173	3-methylphenyl	\$ N N OH		
IIA-174	3-methylphenyl	3 ¹ N		

No.	T-R ²	Q-R ⁴	
ПА-175	3-methylphenyl	ئ ^ا کی ا	
IIA-176	3-methylphenyl		
НА -177	2-fluoro_[[,]]3-chlorophenyl	O N CH3	
ПА-179	2-fluoro_[[,]]3-chlorophenyl	Ş ^N N CH₃	
IIA-180	2-fluoro_[[,]]3-chlorophenyl	3 ¹ 100	
IIA-182	3-chlorophenyl	Ž ^l N→OH	
ПА-183	3-chlorophenyl	Ş. N N N N N N N N N N N N N N N N N N N	
IIA-184	3-chlorophenyl	3 ^N √N~OH	
ПА-187	3-methylphenyl	OH OH	
IIA-188	3 methylphenyl	3 ¹ 20H	

No.	T-R ²	Q-R ⁴	
IIA-190	2-fluoro_[[,]]3-chlorophenyl		
ПА-191	phenyl	Z√N OH	
ПА-192	3,4-dimethoxyphenyl	² ^l n → oH	
IIA-193	3-methylphenyl	3 ¹ NOH	
IIA-194	phenyl		

IV-A

Compounds of Formula IV-A

No.	R	T-R ²	Q-R ⁴
IVA-4	Н	phenyl	CO(pyrrolidin-1-yl)
IVA-5	Me	phenyl	CONHCH ₂ (Ph)
IVA-16	Me	3-Cl-phenyl	CONHCH2(pyridin-4-yl)
IVA-17	Н	5-Cl-phenyl	ZIN OH
IVA-18	Н	5-F-phenyl	CONHCH2(tetrahydrofuran-2-yl)
IVA-19	Me	5,6-F ₂ -phenyl	CO(4-Me-piperidin-1-yl)
IVA-20	Н	4-Cl-phenyl	CONHCH₂(pyrid-4-yl)

No.	R	T-R ²	Q-R ⁴
IVA-21	н	4,5-(OMe) ₂ -phenyl	Z HN N
IVA-22	Me	4,5-Cl₂-phenyl	2 N N O CH ₃

- 19. (Currently amended) A composition comprising a compound according to claim 1[[;]] and a pharmaceutically acceptable carrier.
- 20. (Original) The composition according to claim 19 wherein said compound is formulated in a pharmaceutically acceptable manner for administration to a patient.
- 21. (Currently amended) The composition according to claim 19 further comprising an additional therapeutic agent selected from a chemotherapeutic agent, an anti-inflammatory agent, an immunomodulatory or immunosuppressive agent, a neurotrophic factor, and agent for treating liver disease, an anti-viral agent, an agent for treating a blood disorder, an agent for treating diabetes, or an agent for treating an immunodeficiency disorder.
- 22. (Currently amended) The composition according to claim 20 further comprising an additional therapeutic agent selected from a chemotherapeutic agent, an anti-inflammatory agent, an immunomodulatory or immunosuppressive agent, a neurotrophic factor, and agent for treating liver disease, an anti-viral agent, an agent for treating a blood disorder, an agent for treating diabetes, or an agent for treating an immunodeficiency disorder.
- 23. (Withdrawn) A method of inhibiting ERK or AKT activity in a biological sample, comprising the step of contacting said biological sample with a compound according to claim 1.

24-26. (Canceled)

27. (Withdrawn) A method for treating a disease in a patient comprising the step of administering to said patient a composition according to claim 19, wherein said disease is selected from cancer, stroke, diabetes, hepatomegaly, cardiovascular disease, Alzheimer's disease, cystic fibrosis, viral disease, autoimmune diseases, atherosclerosis, restenosis, psoriasis, allergic disorders, inflammation, neurological disorders, a hormone-related disease, conditions associated with organ transplantation, immunodeficiency disorders, destructive bone disorders, proliferative disorders, infectious diseases, conditions associated with cell death, thrombin-induced platelet aggregation, chronic myelogenous leukemia (CML), liver disease, or pathologic immune conditions involving T cell activation.

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- 28. (Withdrawn) The method according to claim 27 wherein the disease is cancer.
- 29. (Withdrawn) The method according to claim 28 wherein said cancer is selected from breast; ovary; cervix; prostate; testis, genitourinary tract; esophagus; larynx, glioblastoma; neuroblastoma; stomach; skin, keratoacanthoma; lung, epidermoid carcinoma, large cell carcinoma, small cell carcinoma, lung adenocarcinoma; bone; colon, adenoma; pancreas, adenocarcinoma; thyroid, follicular carcinoma, undifferentiated carcinoma, papillary carcinoma; seminoma; melanoma; sarcoma; bladder carcinoma; liver carcinoma and biliary passages; kidney carcinoma; myeloid disorders; lymphoid disorders, Hodgkin's, hairy cells; buccal cavity and pharynx (oral), lip, tongue, mouth, pharynx; small intestine; colon-rectum, large intestine, rectum; brain and central nervous system; or leukemia.
- 30. (Withdrawn) The method according to either of claims 28 or 29 comprising the additional step of administering to said patient a chemotherapeutic agent.
- 31. (Withdrawn) The method according to claim 27 wherein the disease is an autoimmune disease.
- 32. (Withdrawn) The method according to claim 31 wherein said autoimmune disease is selected from psoriasis, SLE Lupus, cystic fibrosis, or conditions associated with organ transplantation.

- 33. (Withdrawn) The method according to claim 27 wherein the disease is a neurological disorder.
- 34. (Withdrawn) The method according to claim 33 wherein said neurological disorder is selected from Alzheimer's Disease, Parkinson's Disease, ALS, epilepsy and seizures, Huntington's disease, or stroke.
- 35. (Withdrawn) The method according to claim 27 wherein the disease is a cardiovascular disease.
- 36. (Withdrawn) The method according to claim 35 wherein said cardiovascular disease is selected from restenosis, cardiomegaly, artherosclerosis, myocardial infarction, or congestive heart failure.
- 37. (Withdrawn) The method according to either of claims 35 or 36 comprising the additional step of administering to said patient a therapeutic agent for treating cardiovascular disease.
- 38. (Withdrawn) The method according to claim 27 wherein the disease is an inflammatory disease.
- 39. (Withdrawn) The method according to claim 38 wherein said inflammatory disease is selected from asthma, rheumatoid arthritis, or atopic dermatitis.
- 40. (Withdrawn) The method according to claim 27 wherein the disease is a liver disease.
- 41. (Withdrawn) The method according to claim 40 wherein said liver disease is selected from hepatomegaly or hepatic ischemia.
 - 42-43. (Canceled)